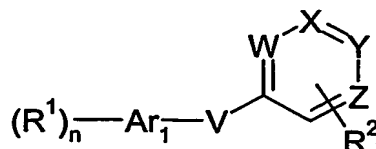


CLAIMS

1. A compound of formula (I):



(I)

wherein

V represents NR⁵, O, S, SO or S(O)₂;

W and X each independently represent CH or N;

Y represents N, CH or C-Ar₂, with the proviso that at least one, but no more than two, of W,

10 X and Y are N;

Z represents CH or C-Ar₂, with the proviso that when Y is N or CH then Z is C-Ar₂, and with the further proviso that when Y is C-Ar₂ then Z is CH;

15 Ar₁ represents a fused 9 or 10 membered heterobicyclic ring system containing one, two, three or four heteroatoms selected from nitrogen, oxygen and sulfur, wherein at least one of the rings in said ring system is aromatic;

Ar₂ represents an aromatic ring selected from phenyl, pyridyl, pyridazinyl, pyrimidinyl and pyrazinyl; which aromatic ring is optionally fused to a phenyl ring, a five-membered heteroaromatic ring containing 1, 2, 3 or 4 heteroatoms selected from O, N and S at most 1 heteroatom being O or S, or a six-membered heteroaromatic ring containing 1, 2 or 3 N atoms; which aromatic ring is unsubstituted or substituted by one, two or three groups selected from halogen, hydroxy, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenylC₁₋₂alkoxy, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy, haloC₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₃₋₇cycloalkyl, C₃₋₇cycloalkoxy, C₃₋₅cycloalkylC₁₋₄alkyl, cyano, nitro, SR⁶, SOR⁶, SO₂R⁶, COR⁶, NR³COR⁶, CONR³R⁴, NR³SO₂R⁶, SO₂NR³R⁴, -(CH₂)_mcarboxy, esterified

25 -(CH₂)_mcarboxy, -(CH₂)_mNR³R⁴, phenyl, naphthyl, a five-membered heteroaromatic ring containing 1, 2, 3 or 4 heteroatoms selected from O, N and S at most 1 heteroatom being O or S and a six-membered heteroaromatic ring containing 1, 2 or 3 N atoms; where two C₁₋₆alkoxy groups are on adjacent atoms they may, together with the atoms to which they are attached, form a 5- or 6-membered partially saturated ring;

30 R¹ represents halogen, hydroxy, oxo, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, haloC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy, haloC₁₋₆alkoxy, hydroxyC₁₋₆alkoxy, C₃₋₇cycloalkyl, C₃₋

cycloalkoxy, C₃₋₅cycloalkylC₁₋₄alkyl, cyano, nitro, SR⁶, SOR⁶, SO₂R⁶, COR⁶, NR³COR⁶, CONR³R⁴, NR³SO₂R⁶, SO₂NR³R⁴, -(CH₂)_mcarboxy, esterified -(CH₂)_mcarboxy or -(CH₂)_mNR³R⁴;

- R² represents hydrogen, halogen, hydroxy, C₁₋₆alkyl, haloC₁₋₆alkyl, C₃₋₇cycloalkyl, C₁₋₆alkoxy, haloC₁₋₆alkoxy, unsubstituted phenyl or phenyl substituted with one or two groups selected from halogen, C₁₋₆alkyl, haloC₁₋₆alkyl, C₃₋₇cycloalkyl, C₁₋₆alkoxy or haloC₁₋₆alkoxy;
- R³ and R⁴ are each independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₃₋₇cycloalkyl or fluoroC₁₋₆alkyl;
- or R³ and R⁴ and the nitrogen atom to which they are attached together form a heteroaliphatic ring of 4 to 7 ring atoms, optionally substituted by one or two groups selected from hydroxy or C₁₋₄alkoxy, which ring may optionally contain as one of the said ring atoms an oxygen or a sulfur atom, S(O), S(O)₂, or NR⁵;
- R⁵ represents hydrogen, C₁₋₄alkyl, hydroxyC₁₋₄alkyl or C₁₋₄alkoxyC₁₋₄alkyl;
- R⁶ represents hydrogen, C₁₋₆alkyl, fluoroC₁₋₆alkyl, C₃₋₇cycloalkyl, unsubstituted phenyl, or phenyl substituted with one or two groups selected from halogen, C₁₋₆alkyl, haloC₁₋₆alkyl, C₃₋₇cycloalkyl, C₁₋₆alkoxy or haloC₁₋₆alkoxy;
- m is either zero or an integer from 1 to 4;
- n is either zero or an integer from 1 to 3;
- or a pharmaceutically acceptable salt, N-oxide or a prodrug thereof.

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2. A compound according to claim 1 in which R¹ is halogen, C₁₋₄alkyl or fluoroC₁₋₄alkyl.
3. A compound according to claim 1 or 2 in which n is one or two.
4. A compound according to claim 1, 2 or 3 in which R² is hydrogen, halogen, C₁₋₄alkyl, C₁₋₄alkoxy or phenyl substituted by C₁₋₄alkyl or fluoroC₁₋₄alkyl.
5. A compound according to any preceding claim in which =W-X=Y- represents =N-CH=CH-, =N-N=CH-, =N-CH=N- or =N-N=C(Ar₂)-.
6. A compound according to any preceding claim in which Ar₁ represents a heterobicyclic ring system selected from isoquinoline, indazole, triazolopyridine, cinnoline, benzothiazole, imidazopyridine, quinoline, tetrahydroisoquinoline or dihydroisoquinoline.
7. A compound according to any preceding claim in which Ar₂ is phenyl or pyridyl which are optionally fused to a phenyl, imidazolyl or thienyl ring, and are unsubstituted or

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substituted by one to three groups independently selected from halogen, cyano, C₁₋₄alkyl, fluoroC₁₋₄alkyl, C₁₋₄alkoxy, fluoroC₁₋₄alkoxy, phenylC₁₋₂alkoxy, piperidine optionally substituted by oxygen, COR⁶ where R⁶ is hydrogen or C₁₋₄alkyl, pyrazole, C₁₋₄alkylcarbonyl, carboxy, C₁₋₆alkylsulphonyl, nitro, phenyl, C₁₋₄alkylthio, hydroxy and -O-CH₂-O-.

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8. A pharmaceutical composition comprising a compound of formula (I) according to any preceding claim, or a pharmaceutically acceptable salt or N-oxide thereof, and a pharmaceutically acceptable excipient.

10 9. A compound of formula (I) according to any one of claims 1 to 7, or a pharmaceutically acceptable salt or N-oxide thereof, for use in a method of treatment of the human or animal body by therapy.

15 10. Use of a compound of formula (I) according to any one of claims 1 to 7, or a pharmaceutically acceptable salt or N-oxide thereof, for use in the manufacture of a medicament for the treatment or prevention of a disease or condition in which pain and/or inflammation predominates.

20 11. A method for the treatment or prevention of a disease or condition in which pain and/or inflammation predominates, which method comprises administration to a patient in need thereof of an effective amount of a compound of formula (I) according to claim 1, or a pharmaceutically acceptable salt or N-oxide thereof.